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SIMULATION OF TRANSDUCER-COUPLANT EFFECTS ON BROADBAND ULTRASONIC SIGNALS

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SIMULATION OF TRANSDUCER-COUPLANT EFFECTS ON

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ABSTRACT

The increasing use of broadband, pulse-echo ultrasonics in nondestructive evaluation of flaws and material properties has generated a need for improved understanding of the way signals are modified by coupled and bonded thin-lay r interfaces associated with transducers. This understanding is most implement when using frequency spectrum analyses for characterizing material properties. In this type of application, signals emanating from material specimens can be strongly influenced by couplant and bond-layers in the acoustic path. Computer sythesized waveforms were used to simulate a range of interface conditions encountered in ultrasonic transducer systems operating in the 20- to 80-MHz regime. The adverse effects of thin-layer multiple reflections associated with various acoustic impedance conditions are demonstrated. The information presented is relevant to ultrasonic transducer design, specimen preparation, and couplant selection.

INTRODUCTION

Ultrasonics for flaw detection and materials characterization is a significant area in nondestructive evaluation (NDE) technology (refs. 1-5). The methodology usually involves broadband transducers in contact with surfaces of test specimens. When frequency spectrum analysis is used for characterizing flaws and material properties, the results can be strongly influenced by couplant and bond-layers associated with the transducer (ref. 1). These thin bond layers and also interconnecting cables can sigmificantly alter the frequency spectra of high-frequency, broadband signals such as those used in making ultrasonic attenuation measurements (refs. 2, 4, 5). For example, spectrum distortions can arise from interference effects due to multiple reflections in thin bond layers. In the case of couplant layers the magnitude of the pressure applied to the ultrasonic probe determines the resultant couplant thickness. Couplant thickness is an important factor in determining the character and acceptability of the signals from a material (refs. 1, 5, 6). The magnitude and nature of signal distortions caused by bond-layer and couplant thickness variations and their related acoustic impedance effects are oftentimes ignored and inadequately understood.

This report treats the effects of thin couplant and bond-layers associated with transducers. Computer simulation methods are used to illustrate the way signals emanating from material specimens can be distorted by thin layers in the acoustic path. Examples are given to demonstrate the adverse effects of thin layers and also coaxial cables. In addition, conditions

under which satisfactory results can be obtained are presented. This paper is believed to be the first attempt to give a systematic account of thin-layer effects as a function of layer thickness and acoustic impedance relative to adjacent materials. The information presented herein is relevant to broadband transducer construction, specimen preparation, and couplant and bond selection and can be an aid in recognizing unacceptable waveforms arising from signal distortions.

APPROACH

The key parameters examined are couplant and bond-layer thickness variations and acoustic impedances of materials commonly occurring in contact ultrasonic involving broadband, buffered probes (refs. 2-5). The frequency range considered is from approximately 20 to 80 MHz, centered at 50 MHz. This range is important in the ultrasonic characterization of the mechanical properties of a variety of materials. It is also a range in which the adverse effects of thin layers become significant. The associated layer thickness are from zero to 50 μm which correspond to the wavelengths involved.

The experimental difficulty of actually varying layer thickness at uniformly spaced intervals from 0 to 50 µm for a number of material combinations is avoided by use of a computer simulation technique. Using this approach, mathematically synthesized waveforms are analyzed by means of a high-speed digital computer and array processing algorithms. The physical acoustics are straightforward, based on the premise of plane elastic waves. The results can be shown to be in excellent agreement with effects that can be observed by direct experimentation, as discussed later.

The transducer-specimen configuration illustrated in figure 1 is taken as a model (refs. 7, 8). As indicated in the figure, the principal material components are an absorber, a piezoelectric element, a buffer, bond layers, a couplant, and the test specimen. The analysis is restricted to consideration of a broadband, ultrasonic pulse signal moving from the specimen into the piezoelement and thence into the adsorber. The buffer serves primarily as a delay line that isolates the piezoelement and specimen. The purpose of the adsorber is to prevent reentry of signals into the piezoelement.

Although the analysis herein treats acoustical reverberations in thin layers, the results are analogous to electronic reflections in coaxial cables used to couple the transducer to a receiver network, as discussed later. In all cases herein, the actual ultrasonic waves in the materials are depicted and referred to in terms of their electrical signal analogs, such as those emitted by the piezoelement in response to a transient pressure wave.

GOVERNING EQUATIONS

A series of configurations, each involving three materials, are treated in accordance with the schedule given in table 1. In each case, the central material is the thin layer of bond or couplant. Transmission of ultrasonic signals through the thin layer is analyzed by using the conventions illustrated schematically in figure 2. As shown in the figure, signal progression is from material [3] through [2] into [1]. (The wave vectors are normal to the interfaces, not oblique as shown for schematic purposes.)

In general, the acoustic impedances of the three materials will differ and hence give rise to the indicated multiple reflections within the thin layer. The signal, E, that emerges in material [1] will tend to be an unresolved composite formed by superposition of the successive thin-layer reflections E_0 through E_N . Once formed, E is unresolvable into its components unless the layer thickness exceeds the mean wavelength of the source signal, S. The spectrum of E will differ from that of S by varying degrees depending on layer thickness, acoustic impedances, and attenuation in the layer.

The transmission, T, and reflection, R, coefficients for the interfaces [1] - [2] and [2] - [3] are given in terms of the acoustic impedances, Z, with dual subscripts indicating direction (ref. 9).

$$T_{21} = \frac{2Z_1}{Z_2 + Z_1}, \ T_{12} = \frac{2Z_2}{Z_2 + Z_1} \tag{1}$$

$$T_{32} = \frac{2Z_2}{Z_3 + Z_2}, T_{23} = \frac{2Z_3}{Z_3 + Z_2}$$
 (2)

$$R_{12} = -R_{21} = \frac{Z_2 - Z_1}{Z_2 + Z_1} \tag{3}$$

$$R_{23} = -R_{32} = \frac{Z_3 - Z_2}{Z_3 + Z_2} \tag{4}$$

The amplitudes of the successive reverberation signals E_0 to E_N are determined by tramission and reflection coefficients and layer thickness, t, and attenuation coefficient, A:

$$E_{0} = ST_{32}T_{21} \exp(-At)$$

$$E_{1} = E_{0}R_{21}R_{23} \exp(-2At)$$

$$E_{2} = E_{0}R_{21}^{2}R_{23}^{2} \exp(-4At)$$

$$E_{N} = E_{0}R_{21}^{N}R_{23}^{N} \exp(-2NAt)$$
(5)

Each successive signal lags the preceding one by the layer "round trip" delay time, d, where d=2t/v, and v is the velocity in the layer. The composite signal E is formed by summing time-domain amplitude waveforms, each displaced by the parenthetically indicated delay:

$$E = E_0 + E_1(-d) + E_2(-2d) + ... + E_N(-Nd)$$
 (6)

This type of summation is readily accomplished by computer processing of waveform arrays.

In the special case where the acoustic impedance of the layer equals or closely approximates that of one contiguous material (i.e., $Z_2 \cong Z_3$ or $Z_2 \cong Z_1$),

where $T_{31} = 2Z_1/(Z_3 + Z_1)$. As layer thickness approaches zero, $E > ST_{31}$.

WAVEFORM SYNTHESIS

The starting waveform, S, is synthesized from ideal amplitude and phase spectra. The procedure is illustrated in figure 3. To assure that the synthesized waveform is authentic, an actual signal is acquired, digitized, and analyzed in polar form by Fourier transformation. By using the real waveform as a model, classical Gaussian amplitude and linear phase spectra are created. Inverse Fourier transformation is used to synthesized a source waveform, S. This procedure assures that any subsequent distortions become evident upon inspection of the composite waveform, E. The synthetic waveform, S, is normalized to 1 volt (minimum to maximum) and thereafter used as a standard source signal for a particular center frequency and bandwidth.

Computer processing of S proceeds with the creation of waveform arrays for E_0 through E_N in accordance with the preceding equations. Economization of computer time requires selection of the smallest permissible number, N, of thin-layer reflections consistent with simulation of actual conditions. Selection of N=10 is satisfactory for the range of materials and conditions investigated herein. This follows from the fact that the amplitude of each successive reflection is diminished exponentially according to the attenuation coefficient of the layer material.

The output composite waveform, E, is synthesized by array addition of corresponding time-domain elements of E_0 through E_N . This corresponds to matrix addition of amplitudes of the N pressure wave components. The composite waveform, E, is subsequently Fourier transformed into polar amplitude and phase spectra and the results are exhibited graphically, as explained in the next section.

RESULTS

The results presented are restricted to a few key material combinations that illustrate pivotal conditions. These materials and acoustical properties are listed in table II. The graphical results and associated data are organized and presented in the order indicated in table I, in seven sets of figures, figures 4 to 10. The first figure in each set summarizes the data associated with the remaining figures. The remaining figures in each set appear in order of increasing layer thickness and show variations in the composite waveform, E, and its amplitude spectrum as layer thickness increases from zero to 40 μm . For all the material combinations examined, the phase spectra remain linear, with no significant change in slope, and exhibit no interesting features with increasing layer thickness. Phase spectra are therefore omitted.

In figures 4 to 10, the OUT/IN amplitude gives the current value of the ratio E/S for each thickness. The RMS (root mean square) energy level is the ratio of the current energy of E relative to its initial value at zero thickness. The variation of the RMS energy level with layer thickness is shown in the summary graphs at the beginning of each set of figures. Spec-

tral skewing is a measure of signal distortion and is computed as percent displacement of the current peak frequency from the nominal (i.e., original) peak frequency associated with the undistorted waveform at zero thickness. The dashed curve in the amplitude spectrum graph is included to show the amount of spectrum distortion that accompanies increasing layer thicknesses.

DISCUSSION

For material combinations having poorly matched acoustic impedances, the least energy transfer (i.e., minimum RMS energy for the composite signal E) occurs at layer thicknesses of 1/4 wavelength (figs. 4(a), 7(a), and 9(a)). Under this condition destructive interference prevails. A secondary maximum follows at layer thicknesses of 1/2 wavelength because of constructive interference. The relative normalized RMS energy levels of these minima and maxima are identical irrespective of the source-signal center frequency. For example, at 20 MHz center frequency, RMS curves are similar to those in figure 4(a) except that they "stretch" to the right, i.e., the 1/4 wavelength minima occur further to the right and initial negative slopes are less.

The adverse effects of multiple reflections in thin layers become apparent by comparing amplitude spectra shown in figures 4, 7, and 9 for 1/2 wavelength layers. For example, figures 4(h), 7(i), and 9(i) illustrate a classical reduction in spectral bandwidth as the result of a "ringing" layer. These figures contrast sharply with the virtually undisturbed broadband spectra obtained with impedance matching (figs. 5, 6, 8, and 10, e.g.). Figures 4, 7, and 9 also illustrate that although RMS energy reduction with increasing layer thickness may appear tolerable, the associated spectral distortions can be quite unacceptable. Certainly, any procedure that relies on spectrum analysis must at least take account of such distortions and avoid them, if possible.

When different materials are combined, there are practical limitations on the ability to control acoustic impedances. By way of compromise, an alternative to perfectly matched impedances would simply require, for example, that

$$Z_1 \le Z_2 < Z_3$$
 or $Z_1 > Z_2 \ge Z_3$

wherein the layer acoustic impedance lies between that of contiguous materials; see figures 5, 6, 8, and 10. In these cases, there is a much smaller loss of signal strength due to reverberations: the RMS level of E drops by less than 5 percent as compared with ~50 percent (fig. 5(a) vs. fig. 4(a)) as layer thickness approaches 1/2 wavelength. Moreover, inteference effects become insignificant relative to attenuation, as predicted by the previous equation for $Z_2 \cong Z_1$ or $Z_2 \cong Z_3$, equation (7). Energy loss due solely to attenuation is indicated by the dashed lines in the first graph of each set (figs. 4 to 10). Examination of figures 5(a), 6(a), 8(a), and 10(a) indicates that for layers with intermediate impedances the composite signal will have only slight or no distortion because it is merely diminished by attenuation in the layer.

Close attention to coupling conditions in contact ultrasonics has been urged by previous investigators (refs. 1, 5, 6). Optimum coupling demands virtually perfect flatness for the buffer-specimen interface and the appli-

cation of substantial force to minimize the couplant layer. Figure 4 shows that in the 20- to 80-MHz range couplant thickness should be less than approximately 1 µm to avoid serious spectral distortions in the case of the typical materials: fused quartz, glycerine, and steel. This requirement is primarily a result of the acoustic mismatch of the glycerine with the contiquous materials. Among the practical alternatives to glycerine that are convenient and safe to use (water, oils, gels, silicones), none have acous-

tic impedances that are significantly different.

It is apparent in figure 5 that an ideal fluid couplant, fluid-X, would have an acoustic impedance close to that of the buffer material (e.g., glass or quartz). It would allow free movement of the transducer over the specimen surface and would relax surface flatness tolerances. Fluid-X also allows the couplant-layer thickness to exceed 10 µm without serious consequences on signal fidelity. Methylene iodide would qualify as fluid-X with an acrylic buffer; see figure 6. In cases where the very low attenuation and ruggedness of fused quartz are preferred, potential candidates for fluid-X are colloidal suspensions of submicrometer particles of metal, metal oxides, or ceramics. Gallium may be useful in restricted applications since it liquefies at ~30°C and readily wets glasses.

The effect of couplant thickness variations shown for synthetic waveforms in figures 4(b) to (1) can be readily verified by applying increasing force to a specimen held against a similar buffered transducer. The source signal can be the first echo from the free back surface of the specimen. As the pressure is increased and the couplant thickness diminishes, a sequence of waveforms on an oscilloscope will duplicate those appearing in the figures. If the buffer and specimen surfaces are sufficiently flat, and the couplant thickness is uniform to within 1 µm, an essentially undistorted waveform will be observed, if the transducer itself is free of internal dis-

torting layers.

As indicated in figures 8 and 10, bond-layers (within the transducer) with intermediate impedances yield good signals over a thickness range of approximately 40 µm. These examples assume that the absorber and bond both consist of tungsten-loaded epoxy. The ideal situation would be to cast and cure the absorber material in place and thus avoid the bond-layer altogether. However, better properties are achieved if the absorber material is formed separately under high-pressure curing (ref. 7). The previous reference also suggests tailoring tungsten-loaded epoxy bond-layers that approach the acoustic impedance of PZT piezoelements (see table 11). To satisfy broadband damping conditions, the bond-layer on either side of the piezoelement should have either slightly higher or slightly lower acoustic impedance (ref. 9). The results presented herein suggest that the bondlayer thickness is not critical under this condition, and therefore, it does not need to be held to a few micrometers.

It can usually be assumed that the ultrasonic receiver, oscilloscope, and associated electronic networks amplify and reproduce signals emitted by the piezoelement in a consistent-manner. However, the coaxial cable that links the transducer to the electronic network can introduce severe distortions in the signal. In this case, the cable is analogous to a thin layer sustaining multiple reflections. Resultant waveform distortions can be shown to be identical to those appearing in figures 4, 7, and 9 as a result of thin-layer reverberations. For coaxial cables having lengths of approximately 1 m, delay times are in excess of 3 nsec, and they are of the same order as delays in thin layers several micrometers thick.

impedance matching of the cable to both the transducer and electronic network. The impedance of the cable should match the 50-ohm terminations conventionally provided in ultrasonic systems. Nevertheless, additional fine actustment may be required and can be provided by adding variable resistors at either end of the cable, as illustrated in figure 11. A "damping' resistor for this purpose is usually shunted across the cable input connector in ultrasonic receivers. A variable, auxiliary damping resistor built into the transducer housing, as in figure 11, greatly enhances signal fidelity. The author has found this auxiliary impedance matching capability indispensible for correcting aberrations peculiar to commercial transducer assemblies. Impedance adjustments at both ends of the cable provide a means to compensate for the effects of electronic and acoustic reverberations.

It is worth becomming familiar with the renegade waveforms shown in the examples given herein. Any waveform having pronounced asymmetry or excess ringdown oscillations should be suspect unless it is recovered from a material sample known to introduce distortions (as with coarse grains, laminations, etc.). Illustrative examples of acceptable and unacceptable waveforms produced by varying the auxiliary damping resistance, and hence the degree of cable impedance matching, appear in figure 12. Any adjustment in coupling, bonding, and cable impedance matching will, of course, change the "system" modulation transfer characteristic. However, these adjustments are discretionary and should be made for convenience in subsequent deconvolutions of signals recovered from specimen materials.

LONCLUSIONS

computer synthesis was used to simulate thin couplant and bond-layer effects associated with broadband ultrasonic transducers. It was shown that these thin layers in the acoustic path can produce distortions in ultrasonic signals and that these distortions become apparent and serious in the frequency regime from approximately 20 to 80 MHz. Selected examples are given to illustrate the potentially adverse effects of thin layers and practical approaches to recognizing and minimizing these effects. The results support the following conclusions:

- l. When couplant or bond-layer acoustic impedances are significantly less than those of both of the contiguous materials joined (as with quartz, glycerine, and steel), the layer thickness should be less than 1 μm to avoid adverse signal distortion effects. This imposes a similar limitation on specimen surface flatness variations, which should be held to fractions of a micrometer in the transducer contact area.
- 2. The preceding toleran's limitation is removed when acoustic impedances of couplant or bond-layers are intermediate between those of the materials joined. In this case, exact matching of the layer and contiguous material acoustic impedance is unnecessary, and the layer thickness can exceed several tens of micrometers. In the cases illustrated, the signal is merely diminished by attenuation in the layer while distortions are minimized or absent.
- 3. Additional corrections to enhance signal fidelity can be made by inserting an auxiliary damping resistor in the transducer housing to complement an input damping resistor in the receiver housing and thereby compensate for adverse reverberation effects introduced by cable or transducer impedance mismatch effects.

The use of computer-simulated experimentation involving synthesized waveforms has clarified questions concerning multiple reverberation effects in thin layers. Additional work using this approach is recommended to study the effects of compound layers within transducers (relative to piezoelements) and within material specimens (with lamellar microstructures).

ACKNOWLEDGEMENT

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REFERENCES

- 1. R. Truell, C. Elbaum, and B. B. Chick, <u>Ultrasonic Methods in Solid State Physics</u>, 1969, pp. 53-151 and 365-368. Academic Press, New York, NY.
- 2. E. P. Papadakis, "Ultrasonic Velocity and Attenuation: Measurement Methods with Scientific and Industrial Applications," <u>Physical Acoustics Principles and Methods</u>, Vol. XII, Chapt. 5, 1976, p. 277.

 Academic Press, New York, NY.
- 3. A. Vary, "Ultrasonic Measurement of Material Properties," Research
 Techniques in Nondestructive Testing, Vol. 1V, Chapt. 5, 1980, p. 160.
 Academic Press, London.
- 4. A. Vary, "Correlations Among Ultrasonic Propagation Factors and Fracture Toughness Properties of Metallic Materials," <u>Materials Evaluation</u>, Vol. 36, June 1978, pp. 55-64.
- 5. A. Vary, "Computer Signal Processing for Ultrasonic Attenuation and Velocity Measurements for Material Property Characterizations," NASA TM-79180 (1979).
- 6. J. H. Williams, Jr., H. Nayeb-Hashemi, and S. S. Lee, "Ultrasonic Attenuation and Velocity in AS/3501-6 Graphite/Epoxy Fiber Composite," NASA CR-3180 (1979).
- 7. K. F. Bainton and M. G. Silk, "Some Factors Which Affect the Performance of Ultrasonic Transducers," <u>British Journal of Non-Destructive Testing</u>, Vol. 22, Jan. 1980, pp. 15-20.
- 8. G. Bradfield, "Ultrasonic Transducers 1. <u>Ultrasonics</u>, Vol. 8, July 1970, pp. 177-189.
- 9. J. Krautkramer and H. Krautkramer, <u>Ultrasonic Testing of Materials</u>, 1969, pp. 20-27 and 100-130. Springer-Verlag, New York, NY.

TABLE 1. - SCHEDULE OF MATERIAL CONFIGURATIONS ANALYZED FOR THIN-LAYER ENERGY TRANSFER AND MULTIPLE-REFLECTION EFFECTS^a

Principal variable ^b	Parametric relation ^c	Mat	Material sequenced			
		[1]	[2]	[3]	figure -	
Couplant thickness	Z ₁ > Z ₂ < Z ₃	Buffer (quartz)	Couplant (glycerine)	Specimen (steel)	4(a)-(1)	
	Z ₁ < Z ₂ < Z ₃	Buffer (quartz)	Louplant (fluid-X)	Specimen (steel)	5(a)-(f)	
·	Z ₁ < Z ₂ < Z ₃	Buffer (acrylic)	Couplant (M-iodide)	Specimen (steel)	6(a)-(f)	
Bond-layer thickness	Z ₁ > Z ₂ < Z ₃	Piezoelement (PZT)	Bond (epoxy)	Buffer (quartz)	7(a)-(1)	
	Z ₁ > Z ₂ > Z ₃	Piezoelement (PZT)	Bond (W-epoxy)	Buffer (quartz)	8(a)-(f)	
Bond-layer thickness	Z ₁ > Z ₂ < Z ₃	Absorber (W-epoxy)	Bond (epoxy)	Piezoelement (PZT)	9(a)-(i)	
	Z ₁ < Z ₂ < Z ₃	Absorber (W-epoxy)	Bond (W-epoxy)	Piezoelement (PZT)	10(a)-(f)	

aResults cover nominal range from 20 to 80 MHz centered at 50 MHz. bThin-layer thickness ranges from zero to 40 µm at 2-µm steps. CAcoustic impedance, Z, is the product of density by (longitudinal) velocity. dMaterial properties and further identification appear in table 11.

TABLE II. - SELECTED MATERIALS, ACOUSTIC PROPERTIES, AND FUNCTIONS

Function	Materiala	Density,b g/cm ³	Velocity ^C cm/µsec	lmpedance ^d g/cm ² µsec
Piezoelectric transduction element	PZT (4 or 5) lead-zirconate niobate ceramic	7.b (7.5-7./)	0.395 (0.38-0.41)	3.0 (2.8-3.2)
Absorber, piezoelement backing	W-epoxy, 40 to 50 percent tungsten powder in epoxy resin	11 & 12 (10-13)	0.21 (0.17-0.24)	2.3 & 2.5 (1.7-3.1)
Adhesive bond	Epoxy resin	1.62	0.26 (0.24-0.28)	0.32 (0.28-0.36)
Buffer, delay	Fused quartz, quartz glass Acrylic resin	1.18	0.595 (0.59-0.60) (0.267-0.27)	1.31 (1.30-1.31) (0.315-0.319)
Couplant	Glycerine Methylene iodide Water (20° 6)	1.26 3.33 1.00	0.192 .098 .148	0.242 .326 .148
Specimen	Mild steel Stainless steel Maraging steel	7.85 7.72 8.03	0.596 .598 .55	4.68 4.62 4.4

aTabulation is limited to materials selected as representative for the purposes of this report in illustrating hin-layer effects.

bProperty values not in parentheses are used for illustrative cases, parenthetical values are quoted to indicate actual range of variation.

CLongitudinal (compressional) ultrasonic wave velocity.

dAcoustic impedance, Z, equals product of density and velocity.

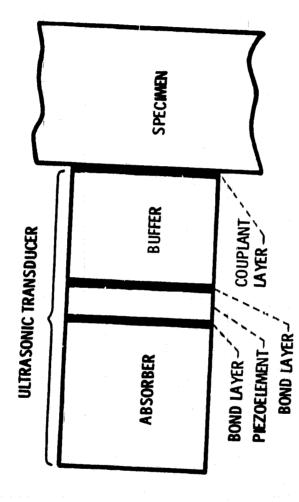


Figure 1. - Diagram of principal components involved in analysis of thin layer reverberations associated with broadband ultrasonic transducers.

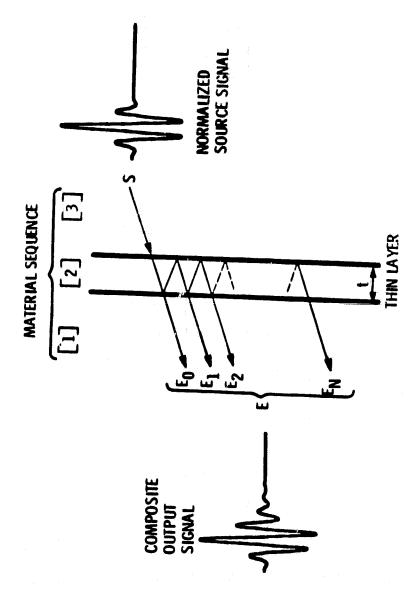


Figure 2. - Diagram of echo system. Ultrasonic signal Sarrising in material [3] emerges in material [1] as a composite signal E resulting from superposition of multiple reflections in thin layer [2].

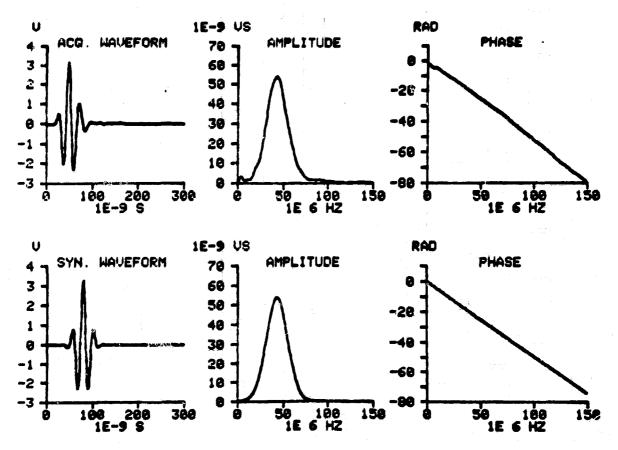


Figure 3. - Time and frequency domain versions of acquired ultrasonic waveform (top graphs) used as model for synthesized waveform (bottom graphs).

THIH LAYER ENERGY TRANSFER AND INTERFERENCE EFFECTS AT 56 MHZ

MATERIAL SYSTEM	DENSITY	UELOCITY	IMPEDANCE
SEQUENCE	(G/CH ³)	(CH/US)	(G/CH ² US)
E13 FUSED DUARTZ (BUFFER) E23 GLYCERINE (COUPLANT) E33 STEEL (SPECIMEN)	2.2 1.26 7.85	.595 .192 .596	1 369 24192 4 6786
THIN LAYER EZJ ATTENUAT	ION COEFFICIENT	- 68 HP/CH (AT	50 MHZ)

COMPOSITE SIGNAL RMS EMERGY (SOLID CURVES) & ATTENUATION (DOTTED CURVE) 1.17 1.1 -1 .9 . 8 .7 .6 .5 .5 6 6 10 12 14 16 40 20 LAYER THICKNESS, HICRONS HAUELEHGTHS IN LAYER X18 (a)

E13 FUSED QUARTZ (BUFFER) E23 GLYCERINE (COUPLANT) E33 STEEL (SPECIMEN)

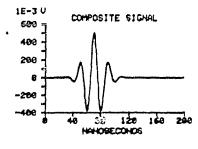
LAYER E23 THICKNESS = 8 MICRON
HAVELENGTHS IN LAYER = 8
AT NOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 8 NANOSEC

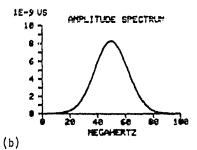
OUT/IN AMPLITUDE = .437237

RHS ENERGY LEUEL = 1

PEAK FREQUENCY = 50 HHZ

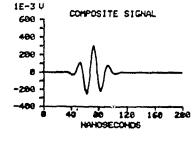
SPECTRAL SKEHING = 0 %





LAYER E23 THICKNESS = 2 MICRON
MAVELENGTHS IN LAYER = 8520633
AT NOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 2.88333 NANOSEC

OUT/IN AMPLITUDE = .272153
RMS ENERGY LEVEL = .618224
PEAK FREDUENCY = 48 MHZ
SPECTRAL SKENING = -4 %



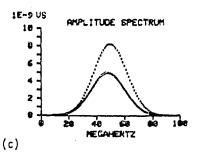


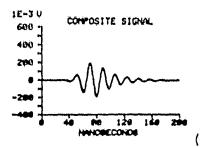
Figure 4. - Degeneration of composite signal and its frequency spectrum as glycerine couplant layer thickness between quartz buffer and steel specimen increases from 0 to 38 micron. Illustration of case where layer impedance is less than that of either contiguous material.

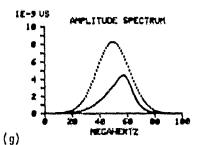
OF MOR QUALTE

EID FUSED OWATZ (BUFFER) EZD GLYCERINE (COUPLANT) EDD STEEL (SPECIMEN)

LAYER E23 THICKNESS = 16 MICRON
MANUELENGTHS IN LAYER = 416667
AT HOMINAL FREQUENCY = 30 MM2
LAYER DELAY = 16 6667 NANOSEC

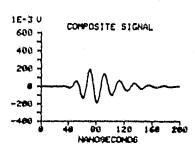
OUT/IN AMPLITUDE = 195958 RMS ENERGY LEVEL = 451947 PEAK FREQUENCY = 57 PHZ SPECTRAL SKENING = 14 %

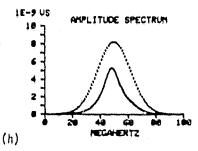




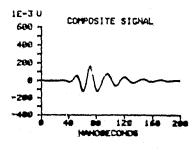
LAYER C23 THICKNESS = 20 HICRON
HAVELENGTHS IN LAYER = 520833
AT NOMINAL FREDUENCY = 50 MHZ
LAYER DELAY = 20 8333 NANOSEC

OUT/IH AMPLITUDE = 189094
RHS ENERGY LEUEL = 480187
PEAK FREQUENCY = 48 1942
SPECTRAL BKEHING = -4 %





LAYER E23 THICKNESS = 24 HICRON HOVELENGTHS IN LAYER = 625 TH 06 HIP FREQUENCY = 50 MH2 LAYER DELAY = 25 NANOSEC OUT/IN AMPLITUDE = .143068 RMS ENERGY LEUEL = .364274 PEAK FREDUENCY = 42 MHZ SPECTRAL SKEWING = -16 %



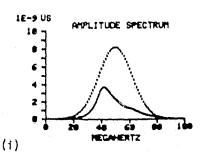
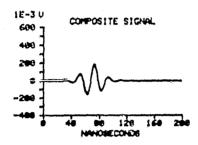
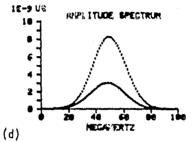


Figure 4. - Continued.

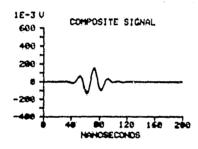
ELD FUSED QUARTZ (BUFFER) EZD GLYCERINE (COUPLANT) E33 STEEL (SPECIMEN)

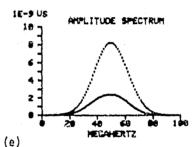
LAYER C23 THICKNESS = 4 MICRON HAVELENGTHS IN LAYER = 184167 AT NOMINAL FREQUENCY = 38 MHZ LAYER DELAY = 4 16667 NANOSEC OUT/IN APPLITUDE = 178061 RMS EMERGY LEVEL = .372811 PEAK FREQUENCY = 49 MHZ SPECTARY SKENING = -2 %





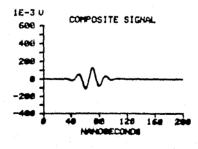
LAYER E23 THICKNESS = 6 MICRON MAUELENGTHS IN LAYER = .13625 AT MOMINAL FREQUENCY = 36 MHZ LAYER DELAY = 6.25 NANOSEC OUT/IN AMPLITUDE = .141845 RMS ENERGY LEVEL = .385478 PEAK FREQUENCY = 56 MHZ SPECTRAL SKEHING = 6 %





LAYER E23 THICKNESS = 10 MICRON
HAVELENGTHS IN LAYER = 260417
AT HOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 10 4167 MANOSEC

OUT/IN AMPLITUDE = .116843 RMS ENERGY LEVEL = .249875 PEAK FREQUENCY = 51 MHZ SPECTRAL SKEHING = 2 %



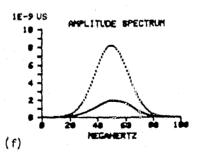
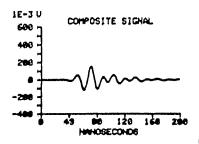
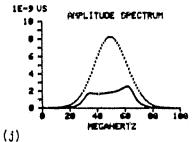


Figure 4. - Continued.

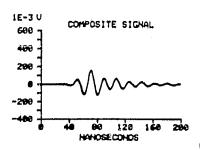
EJD FUSED QUARTZ (BUFFER) EZD GLYCERINE (COUPLANT) EDD STEEL (SPECIMEN)

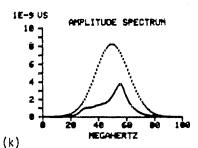
LAYER E23 THICKNESS = 30 MICRON MOVELENGTHS IN LAYER = 78125 AT NOMINAL FREQUENCY = 50 MMZ LAYER DELAY = 31 25 MANOSEC OUT/IN AMPLITUDE = .136343 RMS ENERGY LEVEL = .31467 PEAK FREQUENCY = 62 MHZ SPECTRAL SKEHING = 24 %





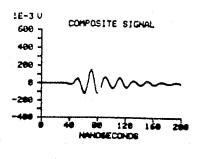
LAYER E23 THICKNESS = 34 MICRON MAUELENGTHS IN LAYER = 885416 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 35 4167 MANOSEC OUT/IN AMPLITUDE = .136162 RMS ENERGY LEVEL = .341745 PEAK FREQUENCY = 56 MHZ SPECTRAL SKEWING = 12 %





LAYER E23 THICKNESS = 38 MICRON
MAUELENGTHS IN LAYER = .989583
AT NOMINAL FREQUENCY = 58 MHZ
LAYER DELAY = 39 5833 MANOSEC

OUT/IN AMPLITUDE = .134407 RMS ENERGY LEVEL = .341605 PEAK FREQUENCY = 50 MHZ SPECTRAL SKEWING = 0 %



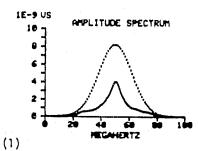


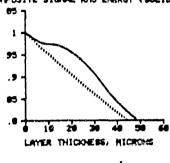
Figure 4. - Concluded.

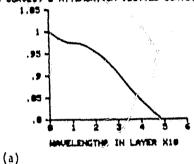
THIN LAYER ENERGY TRANSFER AND INTERFERENCE EFFECTS AT 50 MHZ

MATERIAL SYSTEM	DENSITY	UELOCITY	IMPEDANCE
SEQUENCE E13 FUSED QUARTZ (BUFFER) E23 FLUID-X (COUPLANT) E33 STEEL (SPECIMEN)	(G/CH ³) 2 2 7.85	CCH/US>	(G/CH ² US) 1 389 1 5 4 6786

THIN LAYER EZE ATTENUATION COEFFICIENT - 58 NP/CH (AT 58 MHZ)

COMPOSITE SIGNAL RMS ENERGY (SOLID CURVES) & ATTENDATION (DOTTED CURVE):



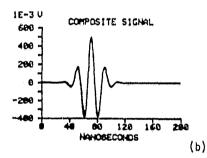


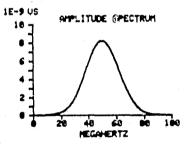
ELD FUSED QUARTZ (BUFFER) E2D FLUID-X (COUPLANT) E3D STEEL (SPECIMEN)

LAYER E23 THICKNESS . 8 MICRON HAVELENGTHS IN LAYER . 0 AT NOMINAL PREQUENCY - 58 7HZ

LAYER DELAY = 8 NAMOSEC

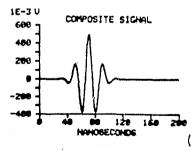
OUT/IN AMPLITUDE = .437237 RMS ENERGY LEVEL - 1 PEAR FREQUENCY = 56 1412 SPECTRAL SKEHING = 0 %





LAYER E23 THICKNESS # 2 MICRON HAUELENGTHS IN LAYER - 02 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY . . 8 NANOSEC

OUT/IN AMPLITUDE = .433913 RMS ENERGY LEUEL # .992232 PEAK FREQUENCY = 50 HHZ SPECTRAL SKEHING = 8 %



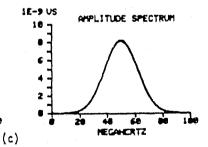
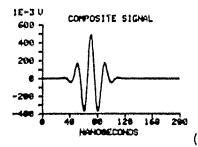
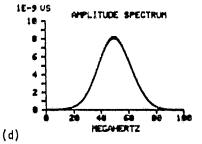


Figure 5. - Variation of composite signal and its frequency spectrum as couplant layer thickness between quartz buffer and steel specimen increases from 0 to 30 micron. Illustration of case where layer impedance is intermediate between that of contiguous materials.

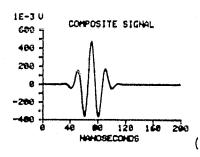
E13 FUSED QUARTZ (BUFFER) E23 FLUID-X (COUPLANT) E33 STEEL (SPECIMEN)

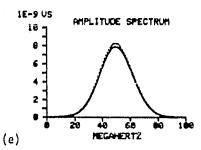
LAYER E23 THICKNESS = 10 MICRON MODELENGTHS IN LAYER = 1 AT NOMINAL FREDUENCY = 50 MHZ LAYER DELAY = 4 NONOSEC DUT/IN AMPLITUDE = 425625 RMS EMERGY LEVEL = .97396 PEAK FREQUENCY = 50 PMZ SPECTRAL SKEWING = 8 %





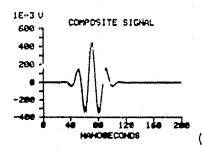
LAYER E23 THICKNESS = 20 MICRON HAVELENGTHS IN LAYER = 2 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 8 MANOSEC OUT/IN AMPLITUDE > .418219
RMS EMERGY LEVEL > .955808
PEAK FREQUENCY = 50 MHZ
EPECTRAL SKEMING = 0 %





LAYER E23 THICKNESS = 39 MICRON
MAUELENGTHS IN LAYER = 3
AT NOMINAL FREDUENCY = 50 MHZ
LAYER DELAY = 12 MANOSEC

OUT/IN AMPLITUDE = .396808 RMS ENERGY LEVEL = .907315 PEAK FREQUENCY = 50 MHZ SPECTRAL SKEWING = 8 %



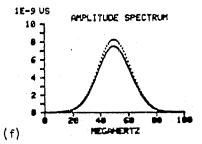
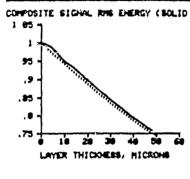


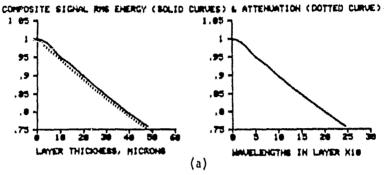
Figure 5. - Concluded.

THIN LAYER ENERGY TRANSFER AND INTERFERENCE EFFECTS AT 58 MIZ

MATERIAL SYSTEM	DENSITY	VELOCITY	IMPEDANCE
SEQUENCE	(G/CH ³)	(chivus)	(C/CH ² US)
EIB ACRYLIC (BUFFER) E2B M-100IDE (COUPLANT) E3B STEEL (GPECIMEN)	1 19 3 33 7. 6 5	27 698 596	.3196 .32634 4.6786

THIN LAYER EZE ATTENUATION COEFFICIENT # 60 MP/CH (AT 50 MHZ)



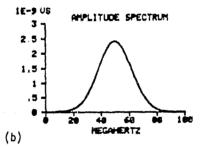


ELID ACRYLIC (BUFFER) EZD METHYLENE TOOTDE (COUPLANT) ESD STEEL (SPECIMEN)

LAYER E23 THICKNESS - 0 HICRON MOVELENGTHS IN LAYER . 0 AT HOMINAL FREQUENCY = 58 MHZ LAYER DELAY . 8 NAMOSEC

1E-3 U COMPOSITE SIGNAL 200 150 100 50 -50 -180 -150 120 NAHORE CONOR

OUTZIN AMPLITUDE = .127511 AMS EMERGY LEVEL = 1 PEAK FREQUENCY = 58 MHZ SPECTRAL SKEHING = 6 %



LAYER C23 THICKNESS = 2 MICRON HQUELENGTHS IN LAYER # .182041 AT NOMINAL FREQUENCY # 50 MHZ LAYER DELAY = 4.88163 HANDSEC

1E-3 U COMPOSITE SIGNAL 150 100 50 . -56 -100 120

OUT/IN AMPLITUDE = .126859 RMS ENERGY LEVEL = .995003 PEAK FREQUENCY = 50 MHZ SPECTRAL SKEHING . 8 %

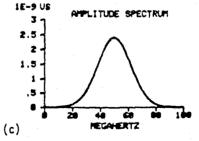
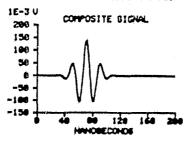


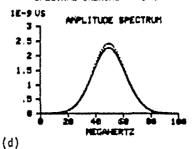
Figure 6. - Variation of composite signal and its frequency spectrum as couplant layer thickness between acrylic buffer and steel specimen increases from 0 to 30 micron. Illustration of case where layer impedance is intermediate between that of contiguous materials.

EID ACRYLIC (BUFFER) E2D METHYLENE 1001DE (COUPLANT) E3D STEEL (SPECIMEN)

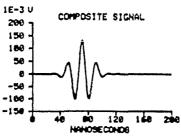
LAYER C23 THICKNESS = 19 MICRON MAVELENGTHS IN LAYER = .518284 AT NOMINAL FREDUENCY = 56 MH2 LAYER DELAY = 20.4882 NANDSEC



OUT/IN AMPLITUDE = .121098 KMS EMERGY LEVEL = .946838 PEAK FREQUENCY = 50 MHZ SPECTRAL SKEHING = 9 %



JAYER E23 THICKNESS = 20 MICRON HAVELENGTHS IN LAYER = 1.02041 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 40 8163 NANOSEC

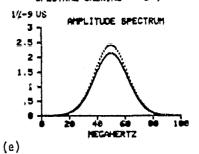


OUT/IN AMPLITUDE = .114268

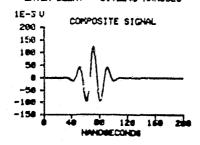
RMS ENERGY LEVEL = .895563

PEAK FREQUENCY = 36 MHZ

SPECTRAL SKEHING = 6 %



LAYER E23 THICKNESS = 38 MICRON
HAVELENGTHS IN LAYER = 1.53861
AT NOMINAL FREQUENCY = 58 MHZ
LAYER DELAY = 61.2245 NANOSEC



OUT/IN AMPLITUDE = .187618 RMS EMERGY LEVEL = .843984 PEAK FREQUENCY = 50 HMZ SPECTRAL DKENING = 0 %

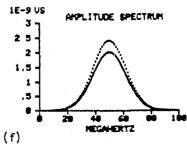


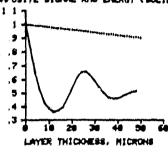
Figure 6. - Concluded.

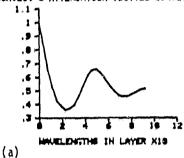
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THIN LAYER ENERGY TRANSFER AND INTERFERENCE EFFECTS AT 50 MHZ

MATERIAL SYSTEM	DEHSITY	VELOCITY	IMPEDANCE
SEQUENCE	(G/CH ³)	(CH/US)	(G/CH ² US)
CID PET (PIEZOELEMENT)	7.6	395 26	3 862
E23 EPOXY (BOHD) E33 FUSED QUARTZ (BUFFER)	1 22	. 595	3172 1 369
THIN LAWS F23 ATTENHATE	ON COMMETCIANT	# 28 MP/CM CAT	56 MIZ)

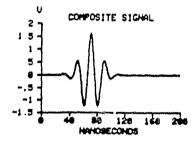
COMPOSITE SIGNAL RMS EMERGY (SOLID CURVES) & ATTENUATION (DOTTED CURVE):

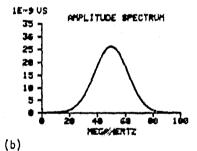




EID PZT (PJEZOELEMENT) E2D EPOXY (BOND) E3D FUSED QUARTZ (BUFFER)

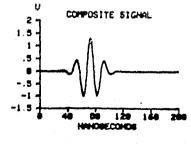
LAYER EGG THICKNESS = 8 MICRON HAVELENGTHS IN LAYER = 8 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 8 MANOSEC OUT/IN AMPLITUDE = 1.39272
RMS ENERGY LEVEL = 1
PEAK FREQUENCY = 56 NHZ
SPECTRAL SKEHING = 16 %





LAYER E23 THICKNESS = 2 MICRON
HAVELENGTHS IN LAYER = .6384613
AT HOMINAL FREQUENCY = 36 PMZ
LAYER DELAY = 1.33846 NANOSEC

OUT/IN AMPLITUDE = 1 15028 RMS ENERGY LEVEL = .820200 PEAK FREQUENCY = 49 HHZ SPECTRAL SKEHING = -2 %



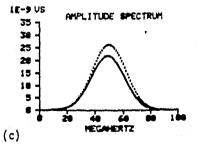
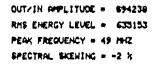
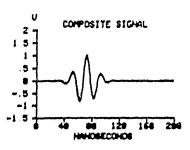


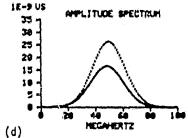
Figure 7. - Degeneration of composite signal and its frequency spectrum as bond layer thickness between PZT piezoelement and quartz buffer increases from 0 to 38 micron. Illustration of case where layer impedance is less than that of either contiguous material.

CID PZY (PIEZOCLEMENT) C2D EPONY (BOND) C3D FUSED QUARTZ (BUFFER)

LAYER E23 THICKNESS = 4 HICRON HOWELENGTHS IN LAYER = 0769231 AT NOMINAL FREQUENCY = 56 MHZ LAYER DELAY = 3 07692 MAHOSEC

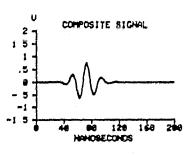


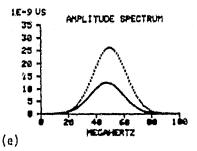




LAYER E23 THICKNESS = 6 MICRON WAVELENGTHS IN LAYER = 115385 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 4 61538 NANOSEC

OUT/IN AMPLITUDE = .677442
RHS EMERGY LEVEL = .476912
PEAK FREQUENCY = 48 HHZ
SPECTRAL SKEHING = -4 %





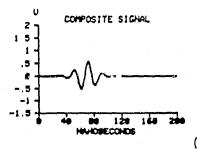
LAYER IZZ THICKNESS = 10 MICRON MAUGLENGTHS IN LAYER = .192306 AT NOMINAL FREQUENCY = 56 MHZ. LAYER DELAY = 7 69231 NANOSEC

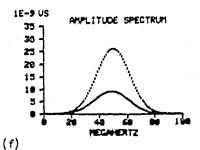
OUT/IN AMPLITUDE = .53392

RMS ENERGY LEVEL = .364316

PEAK FREQUENCY = 49 MHZ

SPECTRAL SKEHING = -2 %



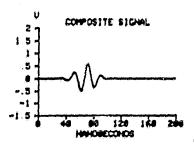


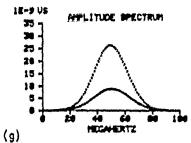
Walderson War

Figure 7. - Continued.

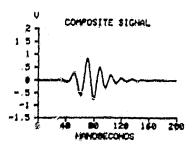
ELD PET (PIEZOCLEMENT) C20 EPOXY (BOND) C30 FUSED GUARTZ (BUFFER)

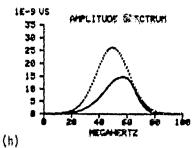
LAYER E23 THICKNESS = 14 HICRON MANUELENGTHS IN LAYER = 269231 HT HOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 10 7692 NONOSEC OUT/IH AMPLITUDE = .526531 RMS ENERGY LEVEL = .352963 PEAK FREQUENCY = 56 MHZ SPECTROL SKENING = .8 %





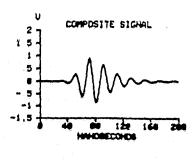
LAYER E23 THICKNESS = 20 MICRON HOVELENGTHS IN LAYER = .304615 AT HOMINAL FREQUENCY = 30 MHZ LAYER DELAY = 15 3046 NANOSEC OUT/IN AMPLITUDE = ,739674 RMS EHERGY LEVEL = .551511 PEAK FREQUENCY = 50 MHZ SPECTROL BKEHING = 16 %





LAYER C23 THICKNESS = 26 MICRON
WWW.ELENGTHS IN LAYER > .5
AT HOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 20 MANOSEC

DUT/IM AMPLITUDE = .8688388 RMS EHERGY LEVEL = .668388 PEAK FREDUENCY = 50 MHZ SPECTRAL SKEWING = 8 %



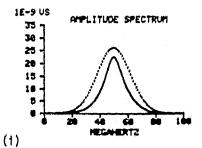
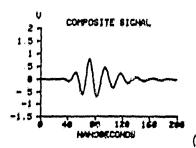
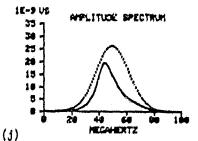


Figure 7. - Continued.

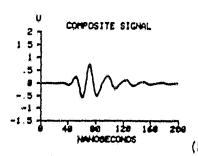
EID PZT (PTEZOELEMENT) EZD EPONY (BOHO) EDD FUSED OUNRYZ (BUFFER)

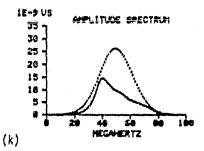
LAYER E23 THICKNESS = 30 MICRON HAWELENGTHS IN LAYER = .576923 AT NOMINAL FREQUENCY = 50 MAZ LAYER DELAY = 23 6769 MANOSEC OUT/IN AMPLITUDE = 72732 RHS EHERGY LEVEL = 501745 PEAK FREQUENCY = 45 MHZ BRECTRAL BREHING = -10 %



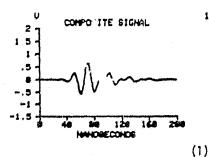


LAYER E23 THICKNESS = 34 MICRON MAUELENGTHS IN LAYER = .653946 AT NOMINAL FREQUENCY = 56 MHZ LAYER DELAY = 26 1538 MANOSEC OUT/IN AMPLITUDE = .648973
RMS ENERGY LEVEL = .48737
PEAK FREQUENCY = 48 NHZ
SPECTRAL SKEHING = -20 %





LAYER E23 THICKNESS = 38 MICRON HAVELENGTHS IN LAYER = .738769 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 29.2308 NANOSEC OUT/IN AMPLITUDE = .638462 RMS ENERGY LEVEL = .451203 PEAK FREQUENCY = 37 HHZ SPECTRAL SKEHING = -26 %



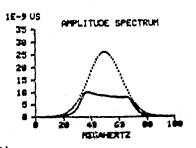
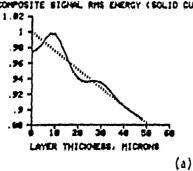


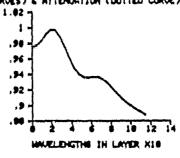
Figure 7. - Concluded.

THIN LAYER ENERGY TRANSFER AND INTERFERENCE REFECTS AT DO MIZ

MATERIAL SYSTEM	DEHSITY	VELOCITY	IMPROMICE
SEQUENCE	(G/Ch ³)	(CH/US)	(G/CH ² US)
EID PZT (PIEZOGLEMENT) EZD W-EMOXY (BOND) EJD FUSED OWARTZ (BUFFER)	7 6 12 2 2	.375 21 595	3 002 2 52 1 349
THIN LAVER COS ATTEMIATE	ON COMPRECIENT	t a 25 NEZCH CAT	56 16(2)

COMPOSITE BIGHAL RHS EMERGY (SOLID CURVES) & ATTEMUATION (DOTTED CURVE)





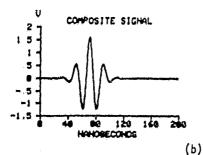
E13 PZT (PIEZOELEMENT) E23 H-EPOXY (BOHD) E33 FUSED QUARTZ (BUFFER)

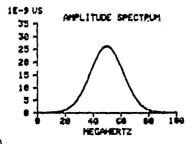
LAYER E23 THICKNESS # 6 MICRON HAVELENGTHS IN LAYER # 6 AT NOMINAL FREQUENCY # 50 MHZ LAYER DELAY # 8 NANOSEC OUT/IN AMPLITUDE = 1 39272

RMS EMERGY LEVEL = 1

PEAK FREQUENCY = 58 MHZ

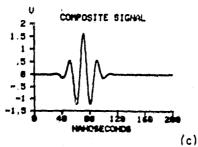
SPECTRAL SKENING = 0 %





LAYER E23 THICKNESS = 2 HICRON MAUELENGTHS IN LAYER = .847619 AT HOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 1.90476 NANOSEC

OUT/IN AMPLITUDE = 1.39315
RMS EMERGY LEVEL = 1.80013
PEAK FREQUENCY = 50 MHZ
SPECTRAL SKEWING = 6 %



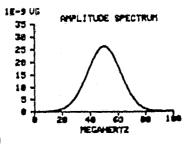
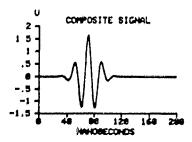


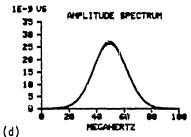
Figure 8. - Variation of composite signal and its frequency spectrum as bond layer thickness between PZT piezoelement and quartz buffer increases from 0 to 30 micron. Illustration of case where layer impedance is intermediate between that of contiguous materials.

E13 PZT (PIEZOELEMENT) E23 H-EPOXY (BONG) E33 FUSED QUARTZ (BUFFER)

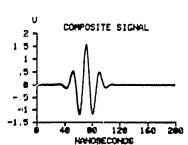
LAYER E23 THICKNESS = 18 HICROH HAVELENGTHS IN LAYER = .238895 AT NOMINAL FREQUENCY = 56 MHZ LAYER DELAY = 9 52381 NANOSE;



OUT/IN AMPLITUSE = 1 43819
RMS ENERGY LEVEL = 1 82563
PEAK FREQUENCY = 58 HHZ
SPECTRAL SKEHING = 8 %



LAYER E23 THICKNESS = 28 MICRON HAWELENGTHS IN LAYER = .47619 AT NOMINAL FREQUENCY = 38 MHZ LAYER DELAY = 19.8476 NANOSEC

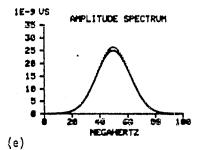


OUT/IN AMPLITUDE # 1.35321

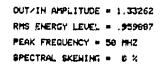
RMS ENERGY LEUEL # .963689

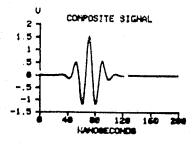
PEAK FREGUENCY # 56 HKZ

SPECTRAL SKEHING # 0 %



LAYER E23 THICKNESS = 30 MICRON MCVELENGTHS IN LAYER = .714286 AT NOMINAL FREQUENCY = 50 MHZ .AYER DELAY = 28 5714 NANOSEC





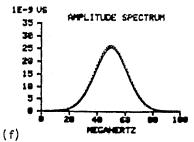
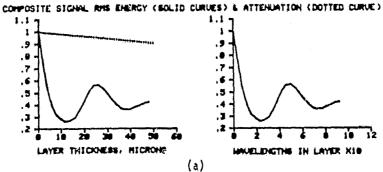
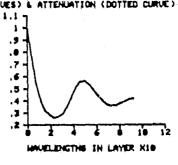


Figure 8. - Concluded.

HATERIAL SYSTEM	DENSITY	VELOCITY	SOMOTH
SEQUENCE	(G/CH ³)	(CH/UE)	(G/CH ² US)
C13 N-EMONY (ABSORBER) C23 EMONY (BOND) C33 MZT (MIEZOELEMENT)	11 1 22 7 6	21 26 395	2 31 3172 3 002
THIN LAYER E23 ATTENUATI	ON COEFFICIENT	- 20 MP/CH CAT	50 M(2)

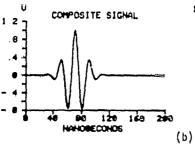


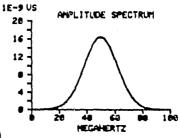


E13 H-EPOXY (ABSORBER) E23 EPOXY (BOND) E33 PZT (PIEZGELEMENT)

LAYER E23 THICKNESS = 8 MICRON HAVELENGTHS IN LAYER . 0 AT NOMINAL FREQUENCY = 58 MHZ LAYER DELAY - 0 NANOSEC

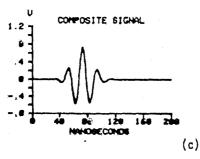
OUT/IN AMPLITUDE = .069729 RMS ENERGY LEVEL = 1 PEAK FREQUENCY = 50 NHZ SPECTRAL SKENING = 0 X





LAYER C23 THICKNESS = 2 MICRON HAUELENGTHS IN LAYER = .8384615 AT NOMINAL FREQUENCY - 58 MHZ LAYER DELAY = 1.53846 NANOSEC

DUT/IN AMPLITUDE = .642242 RHS ENERGY LEVEL # 737171 PEAK FREQUENCY # 49 MHZ SPECTRAL SKEHING = -2 %



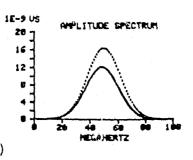
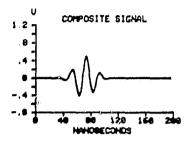
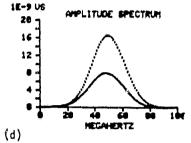


Figure 9. - Degeneration of composite signal and its frequency spectrum as bond layer thickness between tungsten-epoxy absorber and PZT piezoelement increases from 0 to 26 micron. Illustration of case where layer impedance is less than that of either contiguous material.

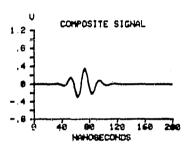
E13 H-EPOXY (ABSORBER) E23 EPOXY (BOND) E33 P2T (PIEZOELEMENT)

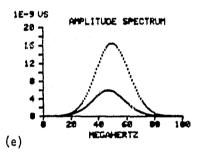
LAYER C23 THICKNESS = 4 MICRON HAUELENGTHS IN LAYER = .0769231 AT NOMINAL FREQUENCY = 50 MMZ LAYER DELAY = 3 07692 NANOSEC OUT/IH AMPLITUDE = .446361 RMS EMERGY LEVEL = .494743 PEAK FREQUENCY = 49 MHZ SPECTRAL SKEHING = -2 %





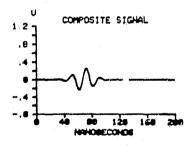
LAYER E23 THICKNESS = 6 MICRON MAUELENGTHS IN LAYER = .115385 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 4.61538 NANOSEC OUT/IN AMPLITUDE = .315181 RMS ENERGY LEVEL = .356727 PEAK FREQUENCY = 47 NHZ SPECTRAL SKEHING = -6 %





LAYER E23 THICKNESS = 10 MICRON
MAUGLENGTHS IN LAYER = .192308
AT NOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 7.69231 NANOSEC

OUT/IN AMPLITUDE = .239925 RMS ENERGY LEVEL = .259472 PEAK FREQUENCY = 48 MHZ SPECTRAL SKEWING = -4 %



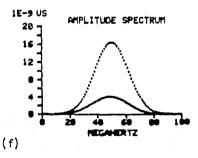


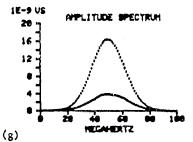
Figure 9. - Continued.

E13 H-EPOXY (ABSORBER) E23 EPOXY (BOND) E33 PZT (PIEZOELEMENT)

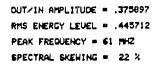
LAYER E23 THICKNESS = 14 MICRON MAUELENGTHS IN LAYER = 269231 AT NOMINAL FREQUENCY = 30 MHZ LAYER DELAY = 10 7692 MANOSEC

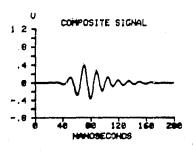


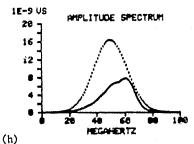
OUT/IN AMPLITUDE = .23442 RMS EMERGY LEVEL = .250415 PEAK FREQUENCY = 49 MHZ SPECTRAL SKEMING = +2 %



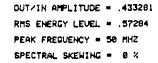
LAYER E23 THICKNESS = 28 MICRON WAVELENGTHS IN LAYER = .384615 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 15.3846 NANOSEC

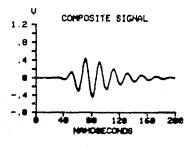






LAYER E23 THICKNESS = 26 MICRON
HAVELENGTHS IN LAYER = .5
AT NOMINAL FREQUENCY = 50 MHZ
LATER DELAY = 20 NANOSEC





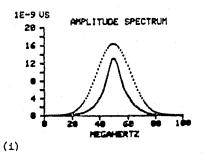
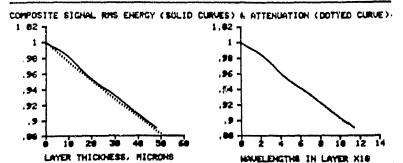


Figure 9. - Concluded.

THIN LAYER ENERGY TRANSFER AND INTERFERENCE EFFECTS AT 50 MHZ

MATERIAL SYSTEM	DEHSITY	VELOCITY	IMPEDANCE
SEQUENCE	(G/CH ³)	(CH/US)	(G/CH ² US)
EID H-EPOXY (ABSORBER) E2D H-EPOXY (BONO) E3D PZT (PIEZOELEMENI)	11 12 7.6	.21 .21 .395	2:31 2:52 3: 06 2
THIN LAYER EST ATTEMAT	ICH COEFFICIENT	= 25 NP/CH (AT	50 MHZ)



E13 H-EPOXY (ABSORBER) E23 H-EPOXY (BOND) E33 PZT (PIEZOELEMENT)

(a)

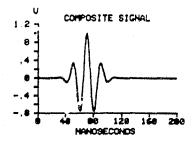
LAYER E23 THICKNESS = 8 MICRON
MAVELENGTHS IN LAYER = 8
AT NOMINAL FREQUENCY = 50 MHZ
LAYER DELAY = 6 NANOSEC

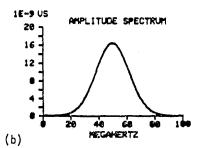
OUT/IN AMPLITUDE = .069729

RMS EMERGY LEVEL = 1

PEAK FREQUENCY = 50 MHZ

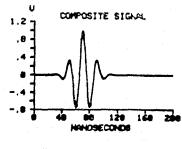
SPECTRAL SKEHING = 0 %





LAYER C23 THICKNESS = 2 MICRON
MAUELENGTHS IN LAYER = 047619
AT NOMINAL FREQUENCY = 30 MHZ
LAYER BELAY = 1.90476 HANOSEC

OUT/IN AMPLITUDE = ,86686
RMS ENERGY LEVEL = .995744
PEAK FREQUENCY = 50 MHZ
SPECTRAL SKEHING = 8 %



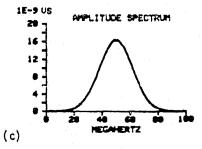
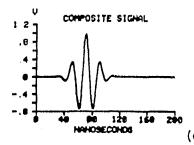
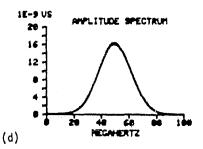


Figure 10. - Variation of composite signal and its frequency spectrum as bond layer thickness between tungstenepoxy absorber and PZT piezoelement increases from 0 to 30 micron. Illustration of case where layer impedance is intermediate between that of contiguous materials.

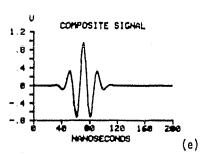
ELD H-EPOXY (ABSORBER) E2D H-EPOXY (BOND) E3D P2T (PIEZOELEMENT)

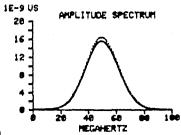
LAYER E23 THICKNESS = 10 MICRON HAVELENGTHS IN LAYER = 238095 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 9 52381 NANOSEC OUT/IN AMPLITUDE = 833395 RMS EMERGY LEVEL = 982899 PEAK FREQUENCY = 50 MHZ SPECTROL SKEHING = 8 %





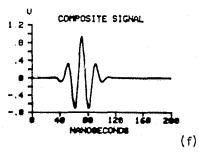
LAYER C23 THICKNESS = 28 MICRON MAUELENGTHS IN LAYER = .47619 AT NOMINAL FREQUENCY = 50 MHZ LAYER DELAY = 19.0476 NANOSEC OUT/IN AMPLITUDE = .829767 RMS ENERGY LEVEL = .932956 PEAK FREQUENCY = 50 MHZ SPECTRAL SKEHING = 0 %





LAYER E23 THICKNESS = 36 MICRON HAUGLENGTHS IN LAYER = .714286 AT NOMINAL FREQUENCY = 56 MHZ LAYER DELAY = 28.5714 NANOSEC

OUT/IN AMPLITUDE = .818181 RMS ENERGY LEVEL = .932112 PEAK FREGUENCY = 50 MHZ SPECTRAL SKEHING = 0 %



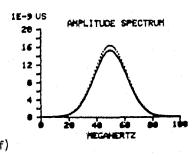


Figure 10. - Concluded.

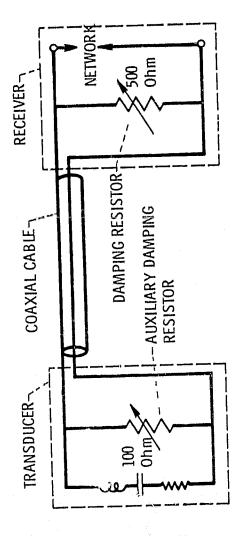


Figure 11. - Diagram showing auxiliary damping resistor included in ultrasonic transducer housing complementing damping resistor in receiver for improved impedance matching. (The piezoelement is supresented by its equivalent reactive components in series, typical values are shown for damping resistors.)

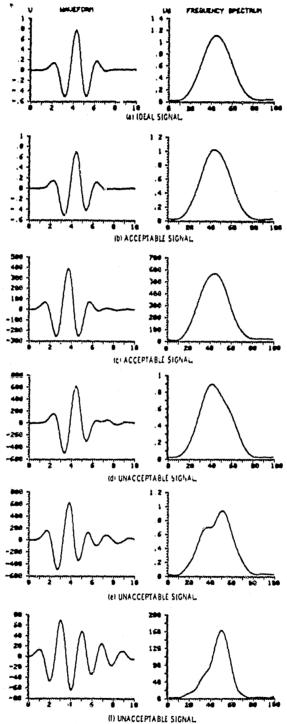


Figure 12. - Waveforms and frequency spectra associated with acceptable and unacceptable signals. Waveform series was generated by increasing electrical impedance mismatch of cable and transducer via the auxiliary damping resistor shown in Fig. 11. Auxiliary damping resistance increases from top to bottom, 2 to 50 0hm, approximately.